Docket No.: 21058/1206459-US2

AMENDMENTS TO THE CLAIMS

Listing of Claims:

This Listing of Claims will replace all prior versions, and listings, of claims in the

application:

1-45 (Cancelled)

46. (New) A method for sequencing nucleic acid, comprising:

a) attaching a template nucleic acid molecule having from about 10 to approximately

100,000 nucleotides in length to a cantilever suitable for detecting a mass dependent property

associated with the cantilever, resulting in forming an attached template nucleic acid; wherein the

attached template nucleic acid is partially double stranded prior to, concurrent with or subsequent

to the attaching of the template nucleic acid;

b) contacting the attached template nucleic acid molecule with at least one type of a

complimentary nucleotide structurally suitable for mass labeling; wherein the complimentary

nucleotide comprises a 3' blocking or protecting group;

c) incubating the attached template nucleic acid molecule and the complimentary nucleotide

under conditions suitable for incorporating the complimentary nucleotide to the attached template

nucleic acid in a position complementary to a nucleotide in the attached template nucleic acid,

wherein the complimentary nucleotide incorporated in the attached template nucleic acid is mass

labeled prior to, concurrent with, or subsequent to the incorporation, yet prior to the addition of a

next complimentary nucleotide; wherein complimentary nucleotides of different types have

different mass labels; and

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d) identifying the complimentary nucleotide incorporated in the attached template nucleic

acid by detecting a change in the mass dependent property associated with the cantilever,

wherein the change is indicative of the incorporation of the complimentary nucleotide in the

attached template nucleic acid.

47. (New) The method of claim 46, wherein the complimentary nucleotide comprises a

chemical structure selected from the group consisting of deoxyadenosine 5' triphosphate (dATP),

deoxythymidine 5' triphosphate (dTTP), deoxyguanosine 5' triphosphate (dGTP) and

deoxycytosine 5' triphosphate (dCTP).

48. (New) The method of claim 46, wherein the complimentary nucleotide comprises a

chemical structure selected from the group consisting of adenosine 5' triphosphate (ATP),

thymidine 5' triphosphate (TTP), guanosine 5' triphosphate (GTP) and cytosine 5' triphosphate

(CTP).

49. (New) The method of claim 46, wherein the change in the mass dependent property of

the structure is determined by detecting deflection and/or resonant frequency shifts in the

cantilever.

50. (New) The method of claim 49, wherein the deflection and/or resonant frequency shift

is detected by optical beam detection, piezoelectric detection, piezoresistance detection or electrical

resistance detection.

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51. (New) The method of claim 46, wherein a single nucleotide polymorphism (SNP) is

identified.

52. (New) The method of claim 46, further comprising iteratively repeating parts b)

through d), wherein for each iteration the attached template is contacted with a different type of

complimentary nucleotide.

53. (New) The method of claim 46, further comprising hybridizing a primer to the attached

template nucleic acid.

54. (New) The method of claim 53, wherein the labeled nucleotides are covalently attached

to the 3' end of the primer by a polymerase.

55. (New) The method of claim 46, wherein the method comprises a plurality of cantilevers,

the cantilevers being arranged in a selected pattern.

56. (New) A method for sequencing nucleic acid, comprising:

a) attaching a template nucleic acid molecule having from about 10 to approximately

100,000 nucleotides in length to a cantilever suitable for detecting a mass dependent property

associated with the cantilever, resulting in forming an attached template nucleic acid; wherein the

attached template nucleic acid is partially double stranded prior to, concurrent with or subsequent

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to the attaching of the template nucleic acid;

b) contacting the attached template nucleic acid molecule with a first type of complimentary

nucleotide structurally suitable for mass labeling, a second type of complimentary nucleotide

structurally suitable for mass labeling, a third type of complimentary nucleotide structurally

suitable for mass labeling, and a fourth type of complimentary nucleotide structurally suitable for

mass labeling; wherein the first, second, third and fourth types of complimentary nucleotides

comprise a 3' blocking or protecting group;

c) incubating the attached template nucleic acid molecule and the first, second, third and

fourth types of complimentary nucleotides under conditions suitable for incorporating the first

type of complimentary nucleotide to the attached template nucleic acid in a position

complementary to a nucleotide in the attached template nucleic acid, wherein the first type of

complimentary nucleotide incorporated in the attached template nucleic acid is mass labeled prior

to, concurrent with, or subsequent to the incorporation, yet prior to the addition of a next

complimentary nucleotide; wherein the first, second, third and fourth types of complimentary

nucleotides have different mass labels; and

d) identifying the first type of complimentary nucleotide incorporated in the attached

template nucleic acid by detecting a change in the mass dependent property associated with the

cantilever, wherein the change is indicative of the incorporation of the first type of complimentary

nucleotide in the attached template nucleic acid.

57. (New) The method of claim 56, wherein one or more of the first, second, third and

37. (New) The method of claim 30, wherein one of more of the first, second, third and

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fourth types of complimentary nucleotides comprise a chemical structure selected from the group

consisting of deoxyadenosine 5' triphosphate (dATP), deoxythymidine 5' triphosphate (dTTP),

deoxyguanosine 5' triphosphate (dGTP) and deoxycytosine 5' triphosphate (dCTP).

58. (New) The method of claim 56, wherein one or more of the first, second, third and

fourth types of complimentary nucleotides comprise a chemical structure selected from the group

consisting of adenosine 5' triphosphate (ATP), thymidine 5' triphosphate (TTP), guanosine 5'

triphosphate (GTP) and cytosine 5' triphosphate (CTP).

59. (New) The method of claim 56, wherein the change in the mass dependent property of

the structure is determined by detecting deflection and/or resonant frequency shifts in the cantilever.

60. (New) The method of claim 59, wherein the deflection and/or resonant frequency shift

is detected by optical beam detection, piezoelectric detection, piezoresistance detection or electrical

resistance detection.

61. (New) The method of claim 56, wherein a single nucleotide polymorphism (SNP) is

identified.

62. (New) The method of claim 56, further comprising iteratively repeating parts b)

through d), wherein for each iteration the attached template is contacted with a different type of

complimentary nucleotide.

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63. (New) The method of claim 56, further comprising hybridizing a primer to the attached

template nucleic acid.

64. (New) The method of claim 63, wherein the labeled nucleotides are covalently attached

to the 3' end of the primer by a polymerase.

65. (New) The method of claim 56, wherein the method comprises a plurality of cantilevers,

the cantilevers being arranged in a selected pattern.

66. (New) A method for sequencing nucleic acid, comprising:

a) attaching a template nucleic acid molecule having from about 10 to approximately

100,000 nucleotides in length to a cantilever suitable for detecting a mass dependent property

associated with the cantilever, resulting in forming an attached template nucleic acid; wherein the

attached template nucleic acid is partially double stranded prior to, concurrent with or subsequent

to the attaching of the template nucleic acid;

b) contacting the attached template nucleic acid molecule of a) with at least one type of a

complimentary nucleotide structurally suitable for mass labeling; wherein the complimentary

nucleotide optionally comprises a 3' blocking or protecting group;

c) incubating the attached template nucleic acid molecule and the complimentary nucleotide

under conditions suitable for incorporating the complimentary nucleotide to the attached template

nucleic acid in a position complementary to a nucleotide in the attached template nucleic acid,

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wherein the complimentary nucleotide incorporated in the attached template nucleic acid is mass

labeled prior to, or concurrent with the incorporation, yet prior to the addition of a next

complimentary nucleotide; wherein complimentary nucleotides of different types have different

mass labels; and

d) identifying the complimentary nucleotide incorporated in the attached template nucleic

acid by detecting a change in the mass dependent property associated with the cantilever,

wherein the change is indicative of the incorporation of the complimentary nucleotide in the

attached template nucleic acid.

67. (New) The method of claim 66, wherein the complimentary nucleotide comprises a

chemical structure selected from the group consisting of deoxyadenosine 5' triphosphate (dATP),

deoxythymidine 5' triphosphate (dTTP), deoxyguanosine 5' triphosphate (dGTP) and

deoxycytosine 5' triphosphate (dCTP).

68. (New) The method of claim 66, wherein the complimentary nucleotide comprises a

chemical structure selected from the group consisting of adenosine 5' triphosphate (ATP),

thymidine 5' triphosphate (TTP), guanosine 5' triphosphate (GTP) and cytosine 5' triphosphate

(CTP).

69. (New) The method of claim 66, wherein the change in the mass dependent property of

the structure is determined by detecting deflection and/or resonant frequency shifts in the cantilever.

70. (New) The method of claim 69, wherein the deflection and/or resonant frequency shift

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is detected by optical beam detection, piezoelectric detection, piezoresistance detection or electrical

resistance detection.

71. (New) The method of claim 66, wherein a single nucleotide polymorphism (SNP) is

identified.

72. (New) The method of claim 66, further comprising iteratively repeating parts b)

through d), wherein for each iteration the attached template is contacted with a different type of

complimentary nucleotide.

73. (New) The method of claim 66, further comprising hybridizing a primer to the attached

template nucleic acid.

74. (New) The method of claim 73, wherein the labeled nucleotides are covalently attached

to the 3' end of the primer by a polymerase.

75. (New) The method of claim 66, wherein the method comprises a plurality of cantilevers,

the cantilevers being arranged in a selected pattern.

76. (New) A method for sequencing nucleic acid, comprising:

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a) attaching a template nucleic acid molecule having from about 10 to approximately

100,000 nucleotides in length to a cantilever suitable for detecting a mass dependent property

associated with the cantilever, resulting in forming an attached template nucleic acid; wherein the

attached template nucleic acid is partially double stranded prior to, concurrent with or subsequent

to the attaching of the template nucleic acid;

b) contacting the attached template nucleic acid molecule with a first type of complimentary

nucleotide structurally suitable for mass labeling, a second type of complimentary nucleotide

structurally suitable for mass labeling, a third type of complimentary nucleotide structurally

suitable for mass labeling, and a fourth type of complimentary nucleotide structurally suitable for

mass labeling; wherein the first, second, third and fourth types of complimentary nucleotides

optionally comprise a 3' blocking or protecting group;

c) incubating the attached template nucleic acid molecule and the first, second, third and

fourth types of complimentary nucleotides under conditions suitable for incorporating the first

type of complimentary nucleotide to the attached template nucleic acid in a position

complementary to a nucleotide in the attached template nucleic acid, wherein the first type of

complimentary nucleotide incorporated in the attached template nucleic acid is mass labeled prior

to, or concurrent with the incorporation, yet prior to the addition of a next complimentary

nucleotide; wherein the first, second, third and fourth types of complimentary nucleotides have

different mass labels; and

d) identifying the first type of complimentary nucleotide incorporated in the attached

template nucleic acid by detecting a change in the mass dependent property associated with the

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cantilever, wherein the change is indicative of the incorporation of the first type of complimentary

nucleotide in the attached template nucleic acid.

77. (New) The method of claim 76, wherein one or more of the first, second, third and

fourth types of complimentary nucleotides comprise a chemical structure selected from the group

consisting of deoxyadenosine 5' triphosphate (dATP), deoxythymidine 5' triphosphate (dTTP),

deoxyguanosine 5' triphosphate (dGTP) and deoxycytosine 5' triphosphate (dCTP).

78. (New) The method of claim 76, wherein one or more of the first, second, third and

fourth types of complimentary nucleotides comprise a chemical structure selected from the group

consisting of adenosine 5' triphosphate (ATP), thymidine 5' triphosphate (TTP), guanosine 5'

triphosphate (GTP) and cytosine 5' triphosphate (CTP).

79. (New) The method of claim 76, wherein the change in the mass dependent property of

the structure is determined by detecting deflection and/or resonant frequency shifts in the cantilever.

80. (New) The method of claim 79, wherein the deflection and/or resonant frequency shift

is detected by optical beam detection, piezoelectric detection, piezoresistance detection or electrical

resistance detection.

81. (New) The method of claim 76, wherein a single nucleotide polymorphism (SNP) is

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identified.

82. (New) The method of claim 76, further comprising iteratively repeating parts b)

through d), wherein for each iteration the attached template is contacted with a different type of

complimentary nucleotide.

83. (New) The method of claim 76, further comprising hybridizing a primer to the attached

template nucleic acid.

84. (New) The method of claim 83, wherein the labeled nucleotides are covalently attached

to the 3' end of the primer by a polymerase.

85. (New) The method of claim 76, wherein the method comprises a plurality of cantilevers,

the cantilevers being arranged in a selected pattern.